## **CLAIMS**

We is claimed is:

- 1. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process comprising the steps of:
  - (a) preparing a silicon carbide fibers solution;
  - (b) preparing a pollen germination medium;
  - (c) preparing a DNA solution;
  - (d) preparing a mixture by mixing said silicon carbide fibers solution and said pollen germination medium with said DNA solution;
  - (e) adding fresh pollens into said mixture to form a paste;
  - (f) vortexing said paste for a time interval of 30-60 seconds;
  - (g) applying said paste for pollination; and
  - (h) selecting for transformants.
- 2. A method for genetic transformation transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 1, wherein said silicon carbide fibers are approximately 0.1- 20  $\mu$ m average diameter and 1 250  $\mu$ m length.
- 3. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 1, wherein the preferred size of said silicon carbide fibers is 1-2  $\mu$ m diameter and  $10-80~\mu$ m length.
- 4. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 1,

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511/<del>1</del>08348.02 02<del>1</del>400/1343/35697.00014 wherein an aqueous solution for silicon carbide fibers is prepared by adding sterile water or solvent to said fibers.

- 5. A method for genetic transformation transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 4, wherein said solution is 5% to 25% aqueous solution.
- 6. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 1, wherein said pollen germination medium is a solution containing about 5% 15% sucrose, 0.01% 1.0% H<sub>3</sub>BO<sub>3</sub>, 0.01% to 1.0% Ca(NO<sub>3</sub>)<sub>2</sub>4H<sub>2</sub>O at pH 5.6.
- 7. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fectandation process according to claim 1, wherein said preferred pollen germination medium is a solution containing about 15% sucrose, 0.018% H<sub>3</sub>BO<sub>3</sub>, 0.04% Ca(NO<sub>3</sub>)<sub>2</sub>4H<sub>2</sub>O at pH 5.6..
- 8. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 1, wherein said DNA is a plasmid DNA.
- 9. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 8, wherein said plasmid DNA is dissolved in a TE solution.
- 10. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 1, wherein said DNA solution is further incubated at about 20 -25<sup>o</sup>C.

- 11. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 1. wherein the selection of a transformate is performed by specific cloned selectable markers having a phenotypic expression or providing resistance to some drugs.
- 12. A method for genetic transformation according to claim 11, wherein said selectable marker having a phenotypic expression is an anthocyanin regulator.
- 13. A method for genetic transformation according to claim 11, wherein said selectable markers providing resistance to some drugs are antibiotics or herbicides.
- 14. A method for genetic transformation according to claim 11, wherein said selectable markers providing resistance to antibiotics is neomycin phosphotransferase gene.
- 15. A method for genetic transformation according to claim 11, wherein said selectable markers providing resistance to antibiotics is kanamycin gene.
- 16. A method for genetic transformation according to claim 11, wherein said selectable markers providing resistance to herbicides is phosphinothricin acetiltransferase gene.
- 17. A method for genetic transformation of any plant species with sexual reproduction based on a pollination-fecundation process according to claim 1, in any plant species with sexual reproduction comprising flowering plants and gymnosperms.
- 18. A method for genetic transformation according to claim 17, wherein said flowering plants are selected from a group consisting of monocots.

- 19. A method for genetic transformation according to claim 18, wherein said monocots is maize.
- 20. A method for genetic transformation according to claim 17, wherein said flowering plants are selected from a group consisting of dicots.
- 21. A method for genetic transformation according to claim 20, wherein said dicots are melon and tomato.
- 22. A method for genetic transformation according to claim 17, wherein said gymnosperms is pine.
- 23. A transgenic maize having an antibiotic kanamycin resistant property prepared by the process of claim 1.
- 24. A transgenic maize having a herbicide bialaphos resistant property prepared by the process of claim 1.
- 25. A transgenic maize having an anthocyanin producing property prepared by the process of claim 1.
- 26. A paste comprising a silicon carbide fiber, a pollen germination medium, and a purified and isolated DNA molecule.
- 27. A paste as recited in claim 26 wherein said silicon carbide fibers having 1-2 μm average
  diameter and 10-80 μm length.
- 28. A paste as recited in claim 26 wherein said silicon carbide fibers is a 5% aqueous solution.

- 29. A paste as recited in claim 26 wherein said pollen germination medium is a solution containing about 15% sucrose, 0.018% H<sub>3</sub>BO<sub>3</sub>, 0.04% Ca(NO<sub>3</sub>)<sub>2</sub>4H<sub>2</sub>O at pH 5.6.
- 30. A paste as recited in claim 26 wherein said DNA is a plasmid DNA.

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